From: Sent: To: Subject:	Russel, Jeffrey Wednesday, Novem STIC-Biotech/Cheml Database Search Re		9.4	
Requester: Jeffrey Russ Art Unit: 1654 Employee Number: 62785 Office Location: REM 3D19 Phone_Number: 571-272-0969 Mailbox Number: REM 3C18			Checked Jer	y-2005
Case serial number 10/810,578 Class / Subclass( NA Earliest Priority NA Format preferred Diskette Search Topic Informate Please search application sequence Geneseq/Uniprot/P Special Instruction	es):  Filing Date: for results:  rmation: h SEQ ID NO:7  nce database PIR. Thank yo	(KEEPPAPPQSP) (pending, publi		
			. 4. 4	NEO 1100 y NO 2 ac Lulyden Lacar (STIO)
Alexand	of Contact: dra Waclawiw II Info. Specialist II 14 30 441  II 3	************  Type of Search  NA# AA#:  S/L: Oligomer:  Encode/Transl:  Structure #:Text:  Inventor: Litigation:	-	Vendors and cost where applicable STN:  DIALOG: QUESTEL/ORBIT: LEXIS/NEXIS: SEQUENCE SYSTEM: WWW/Internet: Other (Specify):

#### => d his ful

L2

(FILE 'HOME' ENTERED AT 09:30:44 ON 03 NOV 2005)

FILE 'REGISTRY' ENTERED AT 09:30:50 ON 03 NOV 2005

FILE 'REGISTRY' ENTERED AT 09:31:23 ON 03 NOV 2005 L1 2 SEA ABB=ON PLU=ON KEEPPAPPQSP/SQSP

FILE 'CAPLUS' ENTERED AT 09:31:43 ON 03 NOV 2005

4 SEA ABB=ON PLU=ON L1

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FILE HOME

### FILE REGISTRY

Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

STRUCTURE FILE UPDATES: 1 NOV 2005 HIGHEST RN 866526-24-1 DICTIONARY FILE UPDATES: 1 NOV 2005 HIGHEST RN 866526-24-1

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TSCA INFORMATION NOW CURRENT THROUGH JULY 14, 2005

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\*\*\*\*\*\*\*\*\*\*\*\*\*\*

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http://www.cas.org/ONLINE/UG/regprops.html

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=> fil reg
FILE 'REGISTRY' ENTERED AT 09:32:25 ON 03 NOV 2005
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New CAS Information Use Policies, enter HELP USAGETERMS for details.

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Please note that search-term pricing does apply when conducting SmartSELECT searches.

Structure search iteration limits have been increased. See HELP SLIMITS for details.

REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

http://www.cas.org/ONLINE/UG/regprops.html

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=> d que l1
L1 2 SEA FILE=REGISTRY ABB=ON PLU=ON KEEPPAPPQSP/SQSP
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=> d l1 sqide3 1-2

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L1 ANSWER 1 OF 2 REGISTRY COPYRIGHT 2005 ACS on STN RN 583028-61-9 REGISTRY
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CN L-Proline, L-lysyl-L-α-glutamyl-L-α-glutamyl-L-prolyl-L-prolyl-L-prolyl-L-glutaminyl-L-seryl- (9CI) (CA INDEX NAME) OTHER NAMES:

CN 10: PN: WO2004052404 SEQID: 10 unclaimed sequence FS PROTEIN SEQUENCE; STEREOSEARCH

SQL 11

unclaimed | SEQID 10

SEQ3 1 Lys-Glu-Glu-Pro-Pro-Ala-Pro-Pro-Gln-Ser-

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11 Pro

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HITS AT: 1-11

\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

MF C52 H81 N13 O18

SR CA

LC STN Files: CA, CAPLUS

DT.CA CAplus document type: Journal; Patent

RL.P Roles from patents: PRP (Properties)

RL.NP Roles from non-patents: BIOL (Biological study); USES (Uses)

Absolute stereochemistry.

PAGE 1-A .

$$H_2N$$
 $(CH_2)_4$ 
 $H_2N$ 
 $(CH_2)_4$ 
 $H_2N$ 
 $(CH_2)_4$ 
 $H_2N$ 
 $(CH_2)_4$ 
 $(CH$ 

PAGE 1-B

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

2 REFERENCES IN FILE CA (1907 TO DATE)

2 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L1 ANSWER 2 OF 2 REGISTRY COPYRIGHT 2005 ACS on STN

RN 348089-26-9 REGISTRY

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L-Proline, L-lysyl-L-\alpha-glutamyl-L-\alpha-glutamyl-L-prolyl-L-prolyl-
    L-alanyl-L-prolyl-L-glutaminyl-O-phosphono-L-seryl- (9CI)
    INDEX NAME)
OTHER NAMES:
    7: PN: US20020147146 SEQID: 7 claimed protein
    7: PN: WO0149709 SEQID: 7 claimed protein
CN
    8: PN: WO2004052404 SEQID: 8 claimed protein
    PROTEIN SEQUENCE; STEREOSEARCH
FS
SQL 11
NTE modified (modifications unspecified)
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                                           description
______
modification
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PATENT ANNOTATIONS (PNTE):
Sequence | Patent
Source
        Reference
Not Given W02001049709
        claimed
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SEQ3
        1 Lys-Glu-Glu-Pro-Pro-Ala-Pro-Pro-Gln-Ser-
          === === === === === === === ===
       11 Pro
          ===
HITS AT:
         1-11
**RELATED SEQUENCES AVAILABLE WITH SEQLINK**
    C52 H82 N13 O21 P
SR
    CA
    STN Files: CA, CAPLUS, USPAT2, USPATFULL
LC
DT.CA Caplus document type: Journal; Patent
      Roles from patents: BIOL (Biological study); PREP (Preparation); PROC
      (Process); PRP (Properties); USES (Uses)
      Roles from non-patents: BIOL (Biological study); USES (Uses)
Absolute stereochemistry.
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- 4 REFERENCES IN FILE CA (1907 TO DATE)
- 4 REFERENCES IN FILE CAPLUS (1907 TO DATE)

#### => fil caplus

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'OBI' IS DEFAULT SEARCH FIELD FOR 'CAPLUS' FILE

=> d que 12

L1 2 SEA FILE=REGISTRY ABB=ON PLU=ON KEEPPAPPQSP/SQSP

L2 4 SEA FILE=CAPLUS ABB=ON PLU=ON L1

=> d .ca 12 1-4

L2 ANSWER 1 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN ACCESSION NUMBER: 2004:513566 CAPLUS Full-text

DOCUMENT NUMBER: 141:47343

TITLE: Glycogen synthase kinase-3 inhibitors

INVENTOR(S): Eldar-Finkelman, Haqit

PATENT ASSIGNEE(S): Tel Aviv University Future Technology Development

L.P., Israel

SOURCE: PCT Int. Appl., 95 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

FAMILY ACC. NUM. COUNT:

English

PATENT INFORMATION:

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APPLICATION NO.
    PATENT NO.
                        KIND
                               DATE
                                                                 DATE
                                           -----
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                               -----
                                          WO 2003-IL1057
    WO 2004052404
                        A2
                               20040624
                                                                  20031211
    WO 2004052404
                        A3
                               20040729
        W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH.
            CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD,
            GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
            LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO,
            NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ,
            TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
        RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ,
            BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE,
            ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK,
            TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD,
                               20040624 CA 2003-2509374 20031211
20050907 EP 2003-777156 20031211
    CA 2509374
                        AA
                        A2
    EP 1569956
        R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
            IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK
PRIORITY APPLN. INFO.:
                                           US 2002-432644P P 20021212
                                                              P 20030627
                                           US 2003-482719P
                                           WO 2003-IL1057
                                                              W 20031211
```

OTHER SOURCE(S): MARPAT 141:47343

ED Entered STN: 25 Jun 2004

Novel conjugates that are capable of inhibiting GSK-3 activity, a process of AB producing same, pharmaceutical compns. including same and methods of using same in the treatment of GSK-3 mediated conditions are disclosed. Methods of treating affective disorders using GSK-3 inhibitors are further disclosed.

ICM A61K047-48 IC

1-10 (Pharmacology) CC

Section cross-reference(s): 2, 63

348089-22-5 **348089-26-9** 348089-28-1 348089-33-8

706785-92-4 706785-93-5

RL: PAC (Pharmacological activity); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(glycogen synthase kinase-3 inhibitors)

149155-45-3 348089-31-6 464875-01-2 **583028-61-9** ·IT

583028-62-0 583028-63-1 708270-79-5 708270-80-8 708270-82-0

708270-83-1 708270-84-2 708270-85-3

RL: PRP (Properties)

(unclaimed sequence; glycogen synthase kinase-3 inhibitors)

ANSWER 2 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN ACCESSION NUMBER: 2003:420432 CAPLUS Full-text

DOCUMENT NUMBER: 139:191139

TITLE: Insulin mimetic action of synthetic phosphorylated

peptide inhibitors of glycogen synthase kinase-3 Plotkin, Batya; Kaidanovich, Oksana; Talior, Ilana;

AUTHOR (S): Eldar-Finkelman, Hagit

CORPORATE SOURCE: Department of Human Genetics and Molecular Medicine,

> Institute of Molecular Medicine, Sackler School of Medicine, Tel Aviv University, Tel Aviv-Jaffa, Israel

SOURCE: Journal of Pharmacology and Experimental Therapeutics

(2003), 305(3), 974-980

CODEN: JPETAB; ISSN: 0022-3565

PUBLISHER: American Society for Pharmacology and Experimental

Therapeutics

Journal

English

DOCUMENT TYPE: LANGUAGE:

ED Entered STN: 02 Jun 2003

Glycogen synthase kinase-3 (GSK-3) was shown to be a key factor in attenuation AB of the cellular action of insulin. The authors speculated that inhibition of GSK-3 might have a potential therapeutic value in treatment of insulin resistance and type 2 diabetes. Here, the authors present a novel class of specific phosphorylated peptide inhibitors of GSK-3, which in sharp contrast to other protein kinase inhibitors that are ATP analogs, are substratecompetitive. The authors show that the GSK-3 peptide inhibitor activated glycogen synthase activity 2.5-fold in human embryonic kidney 293 cells, and increased glucose uptake in primary mouse adipocytes in the absence or presence of insulin compared with cells treated with two resp. peptide controls. In addition, an i.p. administration of GSK-3 peptide inhibitor to normal or insulin-resistant obese C57BL/6J mice, improved their performance on glucose tolerance tests compared with control-treated animals. The authors present here a novel rational strategy for developing specific GSK-3 inhibitors and point toward GSK-3 as a promising therapeutic target in insulin

CC 1-10 (Pharmacology)

Section cross-reference(s): 2

resistance and type-2 diabetes.

IT 158198-86-8 348089-18-9 348089-20-3 348089-22-5 348089-26-9

348089-28-1 348089-31-6 348089-33-8 464875-01-2 583028-59-5

583028-60-8 **583028-61-9** 583028-62-0 583028-63-1

RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(insulin mimetic action of synthetic phosphorylated peptide inhibitors of glycogen synthase kinase-3)

REFERENCE COUNT:

THERE ARE 43 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L2 ANSWER 3 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN ACCESSION NUMBER: 2002:778705 CAPLUS Full-text

DOCUMENT NUMBER:

137:304805

TITLE:

Glycogen synthase kinase-3 inhibitor peptides,

inhibitor design, and therapeutic use

INVENTOR(S):

Eldar-Finkelman, Hagit

PATENT ASSIGNEE(S):

Ramot University Authority for Applied Research &

Industrial Development Ltd., Israel; Tel Aviv University Future Technology Development L.P.

SOURCE:

U.S. Pat. Appl. Publ., 34 pp., Cont.-in-part of Appl.

No. PCT/US01/00123.

CODEN: USXXCO

DOCUMENT TYPE:

Patent English

LANGUAGE:

. 2

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND DATE	APPLICATION NO.	DATE
US 2002147146	A1 20021010	US 2001-951902	20010914
US 6780625	B2 20040824		
WO 2001049709	A1 20010712	WO 2001-US123	20010103
W: AE, AG, AL,	AM, AT, AU, AZ,	BA, BB, BG, BR, BY, BZ,	CA, CH, CN,
CR, CU, CZ,	DE, DK, DM, DZ,	EE, ES, FI, GB, GD, GE,	GH, GM, HR,
HU, ID, IL,	IN, IS, JP, KE,	KG, KP, KR, KZ, LC, LK,	LR, LS, LT,
LU, LV, MA,	MD, MG, MK, MN,	MW, MX, MZ, NO, NZ, PL,	PT, RO, RU,
SD, SE, SG,	SI, SK, SL, TJ,	TM, TR, TT, TZ, UA, UG,	US, UZ, VN,

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            BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
                         A1
                               20040819
                                          US 2004-810578
                                                                  20040329
     US 2004162234
                                                               P 20000103
PRIORITY APPLN. INFO.:
                                           US 2000-174308P
                                                              P 20000522
                                           US 2000-206115P
                                                             A2 20010103
                                           WO 2001-US123
                                           US 2001-951902 . A3 20010914
ED
     Entered STN: 11 Oct 2002
     Peptide inhibitors of glycogen synthase kinase-3 (GSK-3) have an amino acid
AB
     sequence motif of XZXXXS(p)X[S(p) = phosphorylated serine, phosphorylated
     threonine; X = any amino acid; Z = any amino acid except serine or threonine].
     These inhibitors, which are about 7-50 amino acids long, are specific for GSK-
     3 and strongly inhibit the enzyme with an IC50 of about 150 \mu M. Also provided
     are methods of treating biol. conditions mediated by GSK-3 activity, such as
     potentiating insulin signaling in a subject, treating or preventing type 2
     diabetes in a patient, and treating Alzheimer's Disease by administering
     peptide inhibitors. Compns. of these peptide inhibitors and pharmaceutically
     acceptable carriers are also provided, as is a method for identifying
     inhibitors of GSK-3. The invention further relates to a computer-assisted
     method of structure based drug design of GSK-3 inhibitors using a three-
     dimensional structure of a peptide substrate of GSK-3.
IC
     ICM A61K038-17
     ICS C12N009-99
INCL 514012000
     1-12 (Pharmacology)
     Section cross-reference(s): 63
IT
     348089-22-5P 348089-26-9P 348089-28-1P
     RL: PAC (Pharmacological activity); PRP (Properties); SPN (Synthetic
     preparation); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (glycogen synthase kinase-3 inhibitors, inhibitor design, and
        therapeutic use)
                              THERE ARE 80 CITED REFERENCES AVAILABLE FOR THIS
REFERENCE COUNT:
                        80
                              RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
    ANSWER 4 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER: 2001:507722 CAPLUS Full-text
DOCUMENT NUMBER:
                        135:87200
TITLE:
                        Glycogen synthase kinase-3 inhibitors
INVENTOR(S):
                        Eldar-Finkleman, Hagit
PATENT ASSIGNEE(S):
                        Ramot University Authority for Applied Research &
                        Industrial Development Ltd., Israel; McInnis, Patricia
                        Α.
SOURCE:
                        PCT Int. Appl., 49 pp.
                        CODEN: PIXXD2
DOCUMENT TYPE:
                        Patent
                        English
LANGUAGE:
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
     PATENT NO.
                        KIND
                               DATE
                                          APPLICATION NO.
                                                                  DATE
                                          ______
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                               -----
                                          WO 2001-US123
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                               20010712
                         A1
                                                                 20010103
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            HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT,
            LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU,
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SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN,

YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG US 2001-951902 US 2002147146 A1 20021010 20010914 US 6780625 B2 20040824 US 2004162234 Α1 20040819 US 2004-810578 20040329 PRIORITY APPLN. INFO.: US 2000-174308P P 20000103 US 2000-206115P P 20000522 WO 2001-US123 A2 20010103 US 2001-951902 . A3 20010914

ED Entered STN: 13 Jul 2001

The invention is directed to a highly effective and specific peptide inhibitors of glycogen synthase kinase-3 (GSK-3) and useful implications of these peptides. The peptide inhibitors of the invention include therewithin the amino acid motif XZXXXS (p)X, where S(p)=phosphorylated serine or phosphorylated threonine, X=any amino acid, and Z=any amino acid except serine, or threonine. The peptides competitively bind to GSK-3 in vitro with high affinity. Because the amino acid Z in the motif is not phosphorylated, the peptide inhibitor cannot be phosphorylated. Thus, the peptide inhibits the catalytic activity of GSK-3, since the enzyme cannot proceed to phosphorylate other proteins. The peptide inhibitors can be used to diseases mediated by GSK-3 activity such as non-insulin-dependent diabetes mellitus or Alzheimer's disease.

IC ICM C07K007-06

ICS C12N009-12

CC 1-12 (Pharmacology)

Section cross-reference(s): 7

IT 149155-45-3 158198-86-8 348089-18-9 348089-20-3 348089-22-5 348089-24-7 **348089-26-9** 348089-28-1 348089-30-5 348089-31-6 348089-33-8

RL: BAC (Biological activity or effector, except adverse); BPR (Biological process); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); PROC (Process); USES (Uses)

(peptide glycogen synthase kinase-3 inhibitors and their use in treating diseases such as non-insulin-dependent diabetes mellitus and Alzheimer's disease)

REFERENCE COUNT:

THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

# GenCore version 5.1.6 Copyright (c) 1993 - 2005 Compugen Ltd.

OM protein - protein search, using sw model

Run on: November 3, 2005, 08:51:21; Search time 171 Seconds

(without alignments)

32.941 Million cell updates/sec

Title: US-10-810-578-7

Perfect score: 63

Sequence: 1 KEEPPAPPQSP 11

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1612378 seqs, 512079187 residues

Total number of hits satisfying chosen parameters: 1612378

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

왕

Maximum Match 100%

Listing first 150 summaries

Database : UniProt\_03:\*

1: uniprot\_sprot:\*
2: uniprot\_trembl:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	60	95.2	. 213	2	Q7YR98	Q7yr98 bos taurus
` 2	60	95.2	529	1	HSF1_HUMAN	Q00613 homo sapien
3	55	87.3	448	2	Q63717	Q63717 rattus norv
4	52	82.5	525	1	HSF1_MOUSE	P38532 mus musculu
5	49	77.8	231	2	Q6ASV2	Q6asv2 oryza sativ
6	49	77.8	387	2	Q8HXU1	Q8hxul priodontes
7	49	77.8	407	2	Q95KU3	Q95ku3 cabassous u
8	48.5	77.0	562	2	Q8NAF0	Q8naf0 homo sapien
9	48.5	77.0	562	2	Q80VM4	Q80vm4 mus musculu
10	48	76.2	491	1	HSF1_CHICK	P38529 gallus gall
11	47	74.6	94	2	Q8MVN0	Q8mvn0 boltenia vi
12	47	74.6	153	2	Q9NA70	Q9na70 caenorhabdi
13	47	74.6	432	2	Q9ESV6	Q9esv6 rattus norv
14	. 47	74.6	543	1	CH60_BACFO	P81284 bacteroides
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	16	47	74.6	706	2	Q7RXM5		neurospora
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	19	46	73.0	254	1	WAPA_BACST		bacillus st
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	21	46	73.0	282	2	Q9ZR24	Q9zr24	perilla fru
	22	46	73.0	297	1	RPOF_STRAU		streptomyce
	23	46	73.0	303	2	Q9D116	Q9d116	mus musculu
	24	46	73.0	303	2	Q8R3W0	Q8r3w0	mus musculu
	25	46	73.0	319	2	Q8BP27	Q8bp27	mus musculu
	26	46	73.0	319	2	Q9D0D7	Q9d0d7	mus musculu
	27	46	73.0	319	2	Q9D4W4	Q9d4w4	m mus muscu
	28	46	73.0	348	2	Q9CKT8	Q9ckt8	pasteurella
	29	46	73.0	377	2	Q6T3V3		ctenopharyn
	30	46	73.0	471	2	Q8NCK7		homo sapien
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	40		71.4		2	Q8FLQ8		
		45 45		422				corynebacte
	41	45	71.4	433	2	Q7SYN6	_	brachydanio
	42	45	71.4	433	2	Q9PT91		brachydanio
	43	45	71.4	461	2	Q7ZWI6		brachydanio
	44	45	71.4	469	2	Q9V3B8		drosophila
	45	45	71.4	482	2	Q9LZC9		arabidopsis
	46	45	71.4	490	2	Q8Y0Q4		ralstonia s
	47	45	71.4	541	1	PTN5_MOUSE		mus musculu
	48	45	71.4	541	2	Q8CAN0		mus musculu
	49	45	71.4	581	2	Q8 0WU9		mus musculu
ř	50	45	71.4	670	2	Q8GX23	_	arabidopsis
	51	45	71.4	674	2	065672	065672	arabidopsis
	52	45	71.4	681	2	Q9FFW5	Q9ffw5	arabidopsis
	53	45	71.4	738	2	Q82YD4	Q82yd4	streptomyce
	54	45	71.4	757	2	Q7SF38	Q7sf38	neurospora
	55	45	71.4	805	2	092431	092431	bombyx mori
	56	45	71.4	808	1	Y066 NPVAC	P41467	autographa
	57	45	71.4	810	2	Q8B916		rachiplusia
	58	45	71.4	859	2	Q9HFI9		neurospora
	59	45	71.4	978	1	MCR MOUSE		mus musculu
	60	45	71.4	981	1	MCR RAT		rattus norv
	61	45	71.4	1063	2	Q6X2U1		rubella vir
	62	45	71.4	1063	2	Q6X2U3		rubella vir
	63	45	71.4	1135	2	Q812E1		mus musculu
	64	44	69.8	96	2	Q7S8X2		neurospora
	65	44	69.8	128	2	Q852V9		mycobacteri
	66	44	69.8	166	2	Q9UVD0		pneumocysti
	67	44	69.8	299	2	Q89QQ7		bradyrhizob
	68	44	69.8	317	2	Q9D3N6		mus musculu
	69	44	69.8	320	2	Q96BE1		
	70	44	69.8	322	2	Q6PIM1		homo sapien
	70	44	69.8	322				homo sapien
	72	44	69.8	326	2 2	Q8K558		mus musculu
	14	11	09.0	3,40	2	Q89I24	Q09124	bradyrhizob

73	44	69.8	331	2	Q91640	Q9164	0 xenopus lae
74	44	69.8	356	2	Q75HC4	Q75hc	4 oryza sativ
75	44	69.8	368	2	Q6TNR2		2 brachydanio
76	.44	69.8	385	2	Q8HXU7		7 dasypus kap
77	44	69.8	388	2	Q9KIS6	· -	6 brucella ab
78	44	69.8	391	2	Q9RPX5	<del>_</del>	:5 brucella su
79	44	69.8	391	2	Q8FXK6	Q8fxk	6 brucella su
80	44	69.8	393	2	Q6BS91	Q6bs9	1 debaryomyce
81	44	69.8	411	2	Q95KU2		2 dasypus nov
82	44	69.8	417	2	Q20203	· -	3 caenorhabdi
83	44	69.8	471	2	Q6KA29		9 oryza sativ
84	44	69.8	478	2	Q9NXK1		1 homo sapien
85	44	69.8	492	2	Q6C3J1		1 yarrowia li
86	44	69.8	506	2	Q8PDA7	Q8pda	7 xanthomonas
87	44	69.8	510	2	Q8I0Z6	Q8i02	6 caenorhabdi
88	44	69.8	536	2	O84053	08405	3 chlamydia t
89	44	69.8	552	2	Q86YA9		9 homo sapien
90	44	69.8	583	2	Q6YUS6		6 oryza sativ
			590	2			
91	44	69.8			Q91VM4		14 mus musculu
92	44	69.8	593	2	Q8I0Z5		5 caenorhabdi
93	44	69.8	596	2	Q9CVF3		3 mus musculu
94	44	69.8	623	2	Q82HC6	Q82ho	6 streptomyce
95	44	69.8	639	1	GGA1 HUMAN	Q9ujy	5 homo sapien
96	44	69.8	645	2	Q8VDN7	_ <del>_</del>	7 mus musculu
97	44	69.8	656	2	Q61C75		'5 homo sapien
98	44	69.8	712	2	Q8RWX5		5 arabidopsis
99	44	69.8		1			
			722		Z219_HUMAN		4 homo sapien
100	44	69.8	722	2	Q8IYC1		:1 homo sapien
101	44	69.8	726	2	Q66H48	Q66h4	8 rattus norv
102	44	69.8	726	2	Q6IQX8	Q6iq>	8 mus musculu
103	44	69.8	756	2	Q8RP53	Q8rp5	3 streptococc
104	44	69.8	821	2	Q7SC46		6 neurospora
105	44	69.8	839	2	Q69ZV6		6 mus musculu
106	44	69.8	869	2	Q65ZC5		5 caenorhabdi
107	44	69.8	882	2	Q80TZ9	· ·	9 mus musculu
					•	· ·	
108	44	69.8	891	2	Q9VGJ8		8 drosophila
109	44	69.8	937	1	YM92_CAEEL		1 caenorhabdi
110	. 44	69.8	956	2	Q9LJ64	Q91j6	4 arabidopsis
111	44	69.8	957	1	SLK5 MOUSE	Q810h	7 mus musculu
112	44	69.8	958	1	SLK5 HUMAN	09499	1 homo sapien
113	44	69.8	995	2	Q9Y2W4		4 homo sapien
114	44	69.8	1006	2	Q62901		1 rattus norv
115	44	69.8	1012	2	043393		
							3 homo sapien
116	44	69.8	1012	2	075359		9 homo sapien
117	44	69.8	1025	2	Q6NQY8		78 drosophila
118	44	69.8	1026	1	STAU_DROME	P2515	9 drosophila
119	44	69.8	1026	2	Q9V8B9	Q9v81	9 drosophila
120	44	69.8	1114	2	Q6P9L3	Q6p91	3 mus musculu
121	44	69.8	1185	2	Q8VDF4		4 mus musculu
122	44	69.8	1296	2	075046	· · · · · · · · · · · · · · · · · · ·	6 homo sapien
123	44	69.8	1506	2	Q6P6B9		9 homo sapien
124	44	69.8	1566	2	Q9P2R6	<del>-</del>	6 homo sapien
125	44	69.8	1589	2	Q69ZQ7		7 mus musculu
126	44	69.8	1634	2	Q6C908	Q6c90	8 yarrowia li
127	44	69.8	1651	2	Q80TC8	Q80tc	8 mus musculu
128	44	69.8	2321	2	Q6R5R1	Q6r5r	1 meleagrid h
129	44	69.8	2321	2	Q9DGT6		6 meleagrid h
					-	z50	

130	44	69.8	2323	2	Q6HAA3	Q6haa3 meleagrid h
131	44	69.8	2999	2	Q8CHI7	Q8chi7 mus musculu
132	44	69.8	3035	2	Q8CHI8	Q8chi8 mus musculu
133	44	69.8	3084	2	Q8UZ11	Q8uzl1 pseudorabie
134	44	69.8	4340	2	030764	030764 streptomyce
135	43	68.3	118	2	Q6K924	Q6k924 oryza sativ
136	43	68.3	133	2	Q65CJ9	Q65cj9 gallus gall
137	43	68.3	147	1	CHS1_HUMAN	Q9y2v2 homo sapien
138	43	68.3	147	1	CHS1 RAT	Q9wu49 rattus norv
139	43	68.3	163	2	Q9IGT7	Q9igt7 porcine ade
140	43	68.3	167	2	Q7XII6	Q7xii6 oryza sativ
141	43	68.3	167	2	Q73WI5	Q73wi5 mycobacteri
142	43	68.3	178	2	Q9H204	Q9h204 homo sapien
143	43	68.3	193	2	Q9BZJ5	Q9bzj5 homo sapien
144	43	68.3	221	2	Q9X8P1	Q9x8p1 streptomyce
145	43	68.3	222	2	Q9S740	Q9s740 arabidopsis
146	43	68.3	235	2	Q86H42	Q86h42 dictyosteli
147	43	68.3	267	2	Q6P5B3	Q6p5b3 mus musculu
148	43	68.3	315	2	Q66VY4	Q66vy4 gallus gall
149	43	68.3	323	1	PF27 MOUSE	P52875 mus musculu
150	43	68.3	323	2	Q21014	Q21014 caenorhabdi

#### ALIGNMENTS

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07YR98
                 PRELIMINARY;
ID
     Q7YR98
                                    PRT;
                                           213 AA.
AC
     Q7YR98;
DT
     01-OCT-2003 (TrEMBLrel. 25, Created)
     01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT
DT
     01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
DE
     Heat shock transcription factor 1 (Fragment).
GN
     Name=HSF1;
OS
     Bos taurus (Bovine).
OC
     Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC
     Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
     Bovinae; Bos.
OC
OX
     NCBI TaxID=9913;
RN
     [1]
RP
     SEQUENCE FROM N.A.
     PubMed=14667821; DOI=10.1016/S0888-7543(03)00238-6;
RX
RA
     Winter A., Alzinger A., Fries R.;
RT
     "Assessment of the gene content of the chromosomal regions flanking
RT
     bovine DGAT1.";
RL
     Genomics 83:172-180(2004).
DR
     EMBL; AJ518960; CAD58797.1; -.
DR
     GO; GO:0005634; C:nucleus; IEA.
     GO; GO:0003700; F:transcription factor activity; IEA.
DR
     GO; GO:0006457; P:protein folding; IEA.
DR
DR
     GO; GO:0006355; P:regulation of transcription, DNA-dependent; IEA.
DR
     GO; GO:0006986; P:response to unfolded protein; IEA.
DR
     InterPro; IPR000232; HSF DNA bind.
DR
     InterPro; IPR010542; Vert HS TF.
DR
     Pfam; PF00447; HSF DNA-bind; 1.
     Pfam; PF06546; Vert HS TF; 1.
DR
```

RESULT 1

```
KW Heat shock.
    NON_TER
FT
                1
                       1
    NON TER
               213
                    213
FT
              213 AA; 22446 MW; AB675AEA6FF3BA1A CRC64;
    SEQUENCE
SQ
 Query Match
                      95.2%; Score 60; DB 2; Length 213;
 Best Local Similarity 90.9%; Pred. No. 5.3;
 Matches 10; Conservative 1; Mismatches 0; Indels 0; Gaps
                                                                     0;
          1 KEEPPAPPQSP 11
Qу
            1111:1111
Db
        135 KEEPPSPPQSP 145
```

Search completed: November 3, 2005, 09:10:06

Job time : 185 secs

# GenCore version 5.1.6 Copyright (c) 1993 - 2005 Compugen Ltd.

OM protein - protein search, using sw model

Run on: November 3, 2005, 08:54:17; Search time 37 Seconds

(without alignments)

28.605 Million cell updates/sec

Title:

US-10-810-578-7

Perfect score: 63

Sequence:

1 KEEPPAPPQSP 11

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched:

283416 segs, 96216763 residues

Total number of hits satisfying chosen parameters:

283416

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 150 summaries

Database :

PIR\_79:\*

1: pir1:\*

2: pir2:\*

3: pir3:\*

4: pir4:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

### SUMMARIES

		7				•
Result		Query				
No.	Score	Match	Length	DB	ID	Description
			. <b></b>			
1	60	95.2	529	2	A41137	heat shock transcr
2	55	87.3	448	2	S52751	heat shock transcr
3	52	82.5	503	2	A40583	heat shock transcr
4	47	74.6	153	2	T31654	hypothetical prote
5	46	73.0	803	2	G96523	F11A17.8 [imported
6	45	71.4	482	2	T48384	hypothetical prote
7	45	71.4	674	2	T05264	probable serine/th
8	45	71.4	805	2	T41810	AcMNPV orf66 - Bom
9	45	71.4	808	2	C72858	AcOrf-66 protein -
10	45	71.4	929	2	T52517	hypothetical prote
. 11	45	71.4	981	2	A41401	mineralocorticoid
12	44	69.8	417	2	T22024	hypothetical prote
13	44	69.8	460	2	T31587	hypothetical prote

1.4	4.4	60.0	E 2.6	2	H71563	hypothetical prote
14	44	69.8	536	2		_ <del></del>
15	44	69.8	568	2	A34891	Ig heavy chain pre
16	44	69.8	597	2	S40998	hypothetical prote
17	44	69.8	609	2	T14759	hypothetical prote
18	44	69.8	893	2	G88551	protein M01A8.2 [i
19	44	69.8	1006	2	T42731	atrophin-1 related
20	44	69.8	1026	1	A40315	maternal effect pr
21	43	68.3	221	2	T36514	hypothetical prote
22	43	68.3	222	2	H96711	hypothetical prote
23	43	68.3	323	2	T22956	hypothetical prote
24	43	68.3	323	2	A31351	probable transmemb
25	43	68.3	464	2	A47655	spliceosome-associ
26	43	68.3	473	2	T08506	trbI protein - Ent
27	43	68.3	496	2	S26402	homeotic protein H
28	43	68.3	533	2	A56110	
						tyrosine phosphopr
29	43	68.3	684	2	B71379	probable phosphotr
30	43	68.3	700	2	D75508	serine/threonine p
31	43	68.3	802	2	H59434	oligophrenin 1, Rh
32	43	68.3	984	2	A29513	mineralocorticoid
33	42	66.7	148	2	I38881	caudal-type homeot
34	42	66.7	215	2	B75281	peptidyl-tRNA hydr
35	42	66.7	319	2	I38660	melanoma antigen M
36	42	66.7	462	2	E70955	hypothetical prote
37	42	66.7	563	2	T17316	hypothetical prote
38	42	66.7	875	2	T10340	hypothetical prote
39	42	66.7	880	2	T48477	hypothetical prote
40	42	66.7	928	1	RBHU	retinoblastoma-ass
41	42	66.7	1234	2		
					T30160	hypothetical prote
42 -	42	66.7	1293	1	A46350	RNA-directed RNA p
43	42	66.7	1607	2	T03022	MAP kinase kinase
44	41	65.1	183	2	F72697	hypothetical prote
45	41	65.1	226	2	S41032	hypothetical prote
46	41	65.1	250	2	F87483	pentapeptide repea
47	41	65.1	304	1	A60472	uracil-DNA glycosy
48	41	65.1	340	1	QQBES6	UL20 protein precu
49	41	65.1	346	2	T32273	hypothetical prote
50	41	65.1	413	2	H87604	hypothetical prote
51	41	65.1	448	2	I50730	yes-associated pro
52	41	65.1	454	2	A56954	yes-associated pro
53	41	65.1	472	2	B56954	yes-associated pro
54	41	65.1	507	2	S52469	SOX9 protein - mou
55	41	65.1	509	2	A55204	transcription fact
56	41	65.1	575	2	JG0181	X11L2 protein - hu
57	41	65.1	681	2		
					F85062	hypothetical prote
58	41	65.1	731	2	T04455	hypothetical prote
59	41	65.1	757	2	A99561	conserved hypothet
60	41	65.1	759	2	T00875	hypothetical prote
61	41	65.1	845	2	T17291	hypothetical prote
62	41	65.1	902	2	T26775	hypothetical prote
63	41	65.1	920	2	T52426	dynamin-like prote
64	41	65.1	940	2	T00056	hypothetical prote
65	41	65.1	1092	2	H96619	protein T30E16.17
66	41	65.1	1385	2	S61236	major capsid prote
67	41	65.1	1426	2	T30817	homeotic protein C
68	41	65.1	1914	2	T42635	tenascin Y precurs
69	41	65.1	2068	2	A47371	transcription init
70	41	65.1	2649	2	T51023	hypothetical prote
	ملد خد	JJ.1	2047	_	101020	mypochecical proce

### ALIGNMENTS

```
RESULT 1
A41137
heat shock transcription factor 1 - human
C; Species: Homo sapiens (man)
·C;Date: 30-Jun-1992 #sequence revision 30-Jun-1992 #text change 09-Jul-2004
C; Accession: A41137
R; Rabindran, S.K.; Giorgi, G.; Clos, J.; Wu, C.
Proc. Natl. Acad. Sci. U.S.A. 88, 6906-6910, 1991
A; Title: Molecular cloning and expression of a human heat shock factor, HSF1.
A; Reference number: A41137; MUID: 91334376; PMID: 1871105
A; Accession: A41137
A; Molecule type: mRNA
A; Residues: 1-529 < RAB>
A; Cross-references: UNIPROT: Q00613; GB: M64673; NID: g184402; PIDN: AAA52695.1;
PID:g184403
C; Genetics:
A; Gene: GDB: HSF1
A; Cross-references: GDB:128783; OMIM:140580
A; Map position: 8q24.3-8q24.3
C; Superfamily: tomato heat shock transcription factor HSF8; HSF DNA-binding
domain homology
C; Keywords: DNA binding; leucine zipper; transcription regulation
F;18-124/Domain: HSF DNA-binding domain homology <HSF>
                          95.2%; Score 60; DB 2; Length 529;
  Query Match
                          90.9%; Pred. No. 0.88;
  Best Local Similarity
  Matches
            10; Conservative
                                 1; Mismatches
                                                                  0; Gaps
                                                    0; Indels
                                                                              0;
            1 KEEPPAPPQSP 11
Qу
              Db
          298 KEEPPSPPQSP 308
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Search completed: November 3, 2005, 09:10:43
Job time: 44 secs

## GenCore version 5.1.6 Copyright (c) 1993 - 2005 Compugen Ltd.

OM protein - protein search, using sw model

November 3, 2005, 09:10:13; Search time 168 Seconds Run on:

(without alignments)

27.396 Million cell updates/sec

Title: US-10-810-578-7

Perfect score:

Sequence: 1 KEEPPAPPQSP 11

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1867879 seqs, 418409474 residues

Total number of hits satisfying chosen parameters: 1867879

Minimum DB seg length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 150 summaries

Database : Published Applications AA:\*

> 1: /cgn2\_6/ptodata/1/pubpaa/US07\_PUBCOMB.pep:\*

2: /cgn2\_6/ptodata/1/pubpaa/PCT\_NEW\_PUB.pep:\*

3: /cgn2 6/ptodata/1/pubpaa/US06 NEW PUB.pep:\*

/cgn2 6/ptodata/1/pubpaa/US06 PUBCOMB.pep:\* 4:

/cgn2 6/ptodata/1/pubpaa/US07 NEW PUB.pep:\*

/cqn2 6/ptodata/1/pubpaa/PCTUS PUBCOMB.pep:\* 6:

/cgn2\_6/ptodata/1/pubpaa/US08\_NEW\_PUB.pep:\*

8: /cgn2 6/ptodata/1/pubpaa/US08 PUBCOMB.pep:\*

/cgn2 6/ptodata/1/pubpaa/US09A PUBCOMB.pep:\*

10: /cgn2 6/ptodata/1/pubpaa/US09B PUBCOMB.pep:\*

/cgn2\_6/ptodata/1/pubpaa/US09C\_PUBCOMB.pep:\* 11:

12: /cgn2\_6/ptodata/1/pubpaa/US09 NEW PUB.pep:\*

/cgn2 6/ptodata/1/pubpaa/US10A PUBCOMB.pep:\* 13:

/cgn2\_6/ptodata/1/pubpaa/US10B PUBCOMB.pep:\* 14:

/cgn2\_6/ptodata/1/pubpaa/US10C PUBCOMB.pep:\* 15:

16: /cgn2 6/ptodata/1/pubpaa/US10D PUBCOMB.pep:\*

/cgn2\_6/ptodata/1/pubpaa/US10E PUBCOMB.pep:\*

18: /cgn2\_6/ptodata/1/pubpaa/US10 NEW PUB.pep:\*

19: /cgn2 6/ptodata/1/pubpaa/US11A PUBCOMB.pep:\*

20: /cgn2 6/ptodata/1/pubpaa/US11 NEW PUB.pep:\*

/cgn2\_6/ptodata/1/pubpaa/US60\_NEW\_PUB.pep:\*

22: /cgn2 6/ptodata/1/pubpaa/US60 PUBCOMB.pep:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

### SUMMARIES

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US-10-437-963-186486

Sequence 186486,

#### ALIGNMENTS

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; Sequence 7, Application US/09951902
; Patent No. US20020147146A1
; GENERAL INFORMATION:
  APPLICANT: ELDAR-FINKELMAN, Hagit
  TITLE OF INVENTION: GLYCOGEN SYNTHASE KINASE-3 INHIBITORS
  FILE REFERENCE: ELDAR-FINK=1.1B
  CURRENT APPLICATION NUMBER: US/09/951,902
  CURRENT FILING DATE: 2001-09-14
  PRIOR APPLICATION NUMBER: PCT/US01/00123
   PRIOR FILING DATE: 2001-01-03
  PRIOR APPLICATION NUMBER: 60/206,115
  PRIOR FILING DATE: 2000-05-22
  PRIOR APPLICATION NUMBER: 60/174,308
  PRIOR FILING DATE: 2000-01-03
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; Publication No. US20050164324A1
; GENERAL INFORMATION:
  APPLICANT: GYGI, STEVEN P.
  TITLE OF INVENTION: SYSTEMS, METHODS AND KITS FOR CHARACTERIZING
PHOSPHOPROTEOMES
; FILE REFERENCE: 58890(70207)
  CURRENT APPLICATION NUMBER: US/10/862,195
  CURRENT FILING DATE: 2004-06-04
; PRIOR APPLICATION NUMBER: 60/476,010
 PRIOR FILING DATE: 2003-06-04
  NUMBER OF SEQ ID NOS: 2245
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; Publication No. US20030219741A1
; GENERAL INFORMATION:
  APPLICANT: ISOGAI, TAKAO
  APPLICANT: SUGIYAMA, TOMOYASU
  APPLICANT: OTSUKI, TETSUJI
  APPLICANT: WAKAMATSU, AI
  APPLICANT: SATO, HIROYUKI
             ISHII, SHIZUKO
  APPLICANT:
  APPLICANT:
              YAMAMOTO, JUN-ICHI
  APPLICANT: ISONO, YUUKO
  APPLICANT: HIO, YURI
  APPLICANT: OTSUKA, KAORU
  APPLICANT: NAGAI, KEIICHI
             IRIE, RYOTARO
  APPLICANT:
  APPLICANT: TAMECHIKA, ICHIRO
; APPLICANT: SEKI, NAOHIKO
 APPLICANT: YOSHIKAWA, TSUTOMU
  APPLICANT: OTSUKA, MOTOYUKI
  APPLICANT: NAGAHARI, KENJI
  APPLICANT:
              MASUHO, YASUHIKO
  TITLE OF INVENTION: NOVEL FULL-LENGTH CDNA
  FILE REFERENCE: 084335/0160
  CURRENT APPLICATION NUMBER: US/10/094,749
  CURRENT FILING DATE: 2002-03-12
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  PRIOR FILING DATE: 2002-01-24
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  PRIOR FILING DATE: 2001-09-14
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   ORGANISM: Homo sapiens
US-10-094-749-1734
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Search completed: November 3, 2005, 09:24:30 Job time: 174 secs

# GenCore version 5.1.6 Copyright (c) 1993 - 2005 Compugen Ltd.

OM protein - protein search, using sw model

Run on: November 3, 2005, 09:02:33; Search time 41 Seconds

(without alignments)

20.028 Million cell updates/sec

Title: US-10-810-578-7

Perfect score: 63

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Listing first 150 summaries

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; Patent No. 6780625
: GENERAL INFORMATION:
  APPLICANT: ELDAR-FINKELMAN, Hagit
   TITLE OF INVENTION: GLYCOGEN SYNTHASE KINASE-3 INHIBITORS
  FILE REFERENCE: ELDAR-FINK=1.1B
  CURRENT APPLICATION NUMBER: US/09/951,902
  CURRENT FILING DATE: 2001-09-14
; PRIOR APPLICATION NUMBER: PCT/US01/00123
 PRIOR FILING DATE: 2001-01-03
  PRIOR APPLICATION NUMBER: 60/206,115
  PRIOR FILING DATE: 2000-05-22
  PRIOR APPLICATION NUMBER: 60/174,308
 PRIOR FILING DATE: 2000-01-03
  NUMBER OF SEQ ID NOS: 12
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; SEQ ID NO 7
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   OTHER INFORMATION: Synthetic
   NAME/KEY: misc_feature
    LOCATION: (10) .. (10)
    OTHER INFORMATION: Ser residue 10 is phosphorylated.
US-09-951-902-7
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; Sequence 7914, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
  APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
  TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES
THEREOF
; FILE REFERENCE: CL001307
  CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
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; Patent No. 6309820
  GENERAL INFORMATION:
    APPLICANT: SPARKS, Andrew B.
    APPLICANT: HOFFMAN, No. 6309820h
    APPLICANT: KAY, Brian K.
    APPLICANT: FOWLKES, Dana M.
    APPLICANT: McCONNELL, Stephen J.
    TITLE OF INVENTION: POLYPEPTIDES HAVING A FUNCTIONAL
    TITLE OF INVENTION: DOMAIN OF INTEREST AND METHODS OF IDENTIFYING AND
    TITLE OF INVENTION: USING SAME
    NUMBER OF SEQUENCES: 227
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: Pennie & Edmonds LLP
      STREET: 1155 Avenue of the Americas
      CITY: New York
      STATE: New York
      COUNTRY: USA
      ZIP: 10036-2711
    COMPUTER READABLE FORM:
      MEDIUM TYPE: Floppy disk
      COMPUTER: IBM PC compatible
      OPERATING SYSTEM: PC-DOS/MS-DOS
      SOFTWARE: PatentIn Release #1.0, Version #1.30
    CURRENT APPLICATION DATA:
      APPLICATION NUMBER: US/08/630,915A
      FILING DATE: 03-APR-1996
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    ATTORNEY/AGENT INFORMATION:
      NAME: Misrock, S. Leslie
      REGISTRATION NUMBER: 18,872
      REFERENCE/DOCKET NUMBER: 1101-174
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: (212) 790-9090
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      TELEX: 66141 PENNIE
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Search completed: November 3, 2005, 09:11:26

Job time : 43 secs

# GenCore version 5.1.6 Copyright (c) 1993 - 2005 Compugen Ltd.

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November 3, 2005, 08:50:31; Search time 165 Seconds Run on:

(without alignments)

25.784 Million cell updates/sec

Title: US-10-810-578-7

Perfect score: 63

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Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

### SUMMARIES

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7	60	95.2	529	2	AAR13503	Aar13503 HSF. 12/2
8	60	95.2	529	2	AAR24948	Aar24948 Sequence
9	60	95.2	529	2	AAW49093	Aaw49093 Human wil

	10	60	95.2	529	3	AAY55845	Aay55845 Human hea
	11	60	95.2	529	7	ADD48524	Add48524 Human Pro
	12	60	95.2	783	3	AAB22939	Aab22939 GFP-HSF1
	13	60	95.2	783	5	ABG94502	Abg94502 Protease
	14	59	93.7	11	8	ADP47957	Adp47957 GSK-3 inh
	15	57	90.5	11	4	AAG64310	Aag64310 GSK3 pept
	16	5 <i>7</i>	90.5	11	8	ADP47955	Adp47955 Phosphory
	17	57 57	90.5	12	8	ADP47962	
							Adp47962 Phosphory
	18	57	90.5	12	8	ADP47964	Adp47964 GSK-3 inh
	19	55	87.3	448	7	ADD48522	Add48522 Rat Prote
	20	54	85.7	382	3	·AAB58776	Aab58776 Breast an
	21	50	79.4	13	8	ADP47963	Adp47963 N-myristo
	22	48.5	77.0	562	7	ADB65060	Adb65060 Human pro
	23	48.5	77.0	562	8	ADR58973	Adr58973 Human Elk
	24	48.5	77.0	562	8	ADR58971	Adr58971 Human Elk
	25	47	74.6	432	8	ADP44550	Adp44550 Human gly
	26	46	73.0	94	4	AAU31792	Aau31792 Novel hum
	27	46	73.0	282	8	ADJ50141	Adj50141 Oil-assoc
	28	46	73.0	295	7	ADC87211	Adc87211 Human GPC
	. 29	46	73.0	308	5	ABB80594	Abb80594 Human sbg
	30	46	73.0	308	5	AAE21157	Aae21157 Human TRI
i	31	46	73.0	375	4	AAB88570	Aab88570 Human hyd
	32	46	73.0	447			
					5	ABB80595	Abb80595 Human sbg
	33	46	73.0	447	5	ADI16488	Adi16488 Human NOV
	34	46	73.0	447	8	ADN42144	Adn42144 Human nov
	35	46	73.0	471	4	AAM93737	Aam93737 Human pol
	36	46	73.0	471	8	ADL31672	Adl31672 Human pro
	37	46	73.0	472	5	ADI16494	Adi16494 Human NOV
	38	46	73.0	472	8	ADN42150	Adn42150 Human nov
	39	46	73.0	572	2	AAW31855	Aaw31855 Mycobacte
	40	46	73.0	724	5	ABB83924	Abb83924 Arabidops
	41	46	73.0	763	2	AAW31852	Aaw31852 Mycobacte
_	42	45	71.4	126	4	AAO00517	Aao00517 Human pol
	43	45	71.4	195	3	AAB42025	Aab42025 Human ORF
	44	45	71.4	389	5	ABP65964	Abp65964 Bifidobac
	45	45	71.4	469	4	ABB68765	
	46	45	71.4	541	5	ABB57066	Abb57066 Mouse isc
	47	45	71.4	559	8	ADJ67952	
	48	45	71.4	559	8		Adj67952 G. stearo
	49				_	ADJ68164	Adj68164 G. stearo
		45	71.4	559	8	ADK01242	Adk01242 DNA polym
	50	45	71.4	559	8 .	ADJ79461	Adj79461 G. stearo
	51	45	71.4	559	8	ADJ84901	Adj84901 B. steart
	52	45	71.4	559	8	ADM77689	Adm77689 DNA polym
	53	45	71.4	559	8	ADM66356	Adm66356 G. stearo
	54	45	71.4	559	8	ADO04409	Ado04409 B. steart
	55	45	71.4	559	8	ADP82486	Adp82486 B. stearo
	56	45	71.4	674	4	AAB74208	Aab74208 Protein e
	57	45	71.4	674	5	ABB93235	Abb93235 Herbicida
	58	45	71.4	674	7	ABR82941	Abr82941 Arabidops
	59	45	71.4	681	5	ABB93650	Abb93650 Herbicida
	60	45	71.4	830	4	ABB64695	Abb64695 Drosophil
	61	45	71.4	907	8	ADS21481	Ads21481 Bacterial
	62	44	69.8	59	4	ABG27595	
	63	44	69.8	65	8	AB056401	Abg27595 Novel hum
	64	44	69.8	128			Abo56401 Human gen
	65				7	ABO64369	Abo64369 Klebsiell
	66	44	69.8	135	5	AAE22219	Aae22219 Rodent to
	00	44	69.8	185	4	AAU15990	Aau15990 Human nov

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#### ALIGNMENTS

```
RESULT 1
AAG64309
ID
     AAG64309 standard; peptide; 11 AA.
XX
AC
     AAG64309;
XX
DT
     21-SEP-2001 (first entry)
XX
DE
     GSK3 peptide inhibitor #2.
XX
KW
     Antidiabetic; antidepressant; peptide inhibitor; manic depression;
KW
     glycogen synthase kinase-3; GSK3;
     non-insulin dependent diabetes mellitus; Alzheimer's disease.
KW
XX
OS
     Synthetic.
XX
FH
                     Location/Qualifiers
     Key
FT
     Modified-site
FT
                     /label= Phosphoserine
XX
PN
     WO200149709-A1.
XX
PD
     12-JUL-2001.
XX 、
     03-JAN-2001; 2001WO-US000123.
PF
XX
     03-JAN-2000; 2000US-0174308P.
PR
PR
     22-MAY-2000; 2000US-0206115P.
XX
PΑ
     (UYRA-) UNIV RAMOT APPLIED RES & IND DEV LTD.
     (MCIN/) MCINNIS P A.
PΑ
XX
PΙ
     Eldar-Finkleman H;
XX
DR
     WPI; 2001-451785/48.
XX
PT
     Peptide inhibitors of glycogen synthase kinase-3, useful for preventing
PT
     and treating non-insulin dependent diabetes mellitus, Alzheimer's disease
PΤ
     and manic depression.
XX
PS
     Example 1; Page 30; 49pp; English.
XX
CC
     The present invention relates to peptide inhibitors of glycogen synthase
     kinase-3 (GSK3). The peptide inhibitors may be used to inhibit the
CC
CC
     catalytic activity of GSK3 and treat diseases related to inappropriate
     expression and activity of GSK3, e.g. non-insulin dependent (NID)
CC
CÇ
     diabetes mellitus, NID Alzheimer's disease or NID manic depression. In
CC
     particular, the peptide inhibitor is administered prophylactically to
CC
     prevent NID diabetes mellitus by potentiating insulin signalling in type-
CC
     2 diabetics. The present sequence is one such peptide inhibitor for GSK3
XX
SO
     Sequence 11 AA;
```

100.0%; Score 63; DB 4; Length 11;

Query Match

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Best Local Similarity
                          100.0%; Pred. No. 0.13;
            11; Conservative
                                 0; Mismatches
                                                   0;
                                                       Indels
                                                                  0; Gaps
            1 KEEPPAPPQSP 11
Qу
              Db
            1 KEEPPAPPOSP 11
RESULT 2
ADP47956
ID
     ADP47956 standard; peptide; 11 AA.
XX
AC
     ADP47956;
XX
     09-SEP-2004 (first entry)
DT
XX
DE
     GSK-3 inhibitor peptide, seq id 10.
XX
KW
     Anorectic; antidiabetic; neuroleptic; antidepressant; cerebroprotective;
KW
     tranquiliser; vulnerary; neuroprotective; nootropic; anticonvulsant;
KW
     antiparkinsonian; glycogen synthase kinase-3; GSK-3; insulin signaling;
KW
     beta-catenin; obesity; non-insulin dependent diabetes mellitus;
KW
     neurodegenerative disease; psychotic disease; unipolar disorder;
     bipolar disorder; manic depression; ischaemia; stroke; brain injury;
KW
KW
     bacterial infection; Alzheimer's disease; Huntington's disease;
KW
     Parkinson's disease; AIDS associated dementia;
KW
     amyotrophic lateral sclerosis; AL; multiple sclerosis; schizophrenia.
XX
OS
     Unidentified.
XX
PN
     WO2004052404-A2.
XX
PD
     24-JUN-2004.
XX
ΡF
     11-DEC-2003; 2003WO-IL001057.
XX
PR
     12-DEC-2002; 2002US-0432644P.
PR
     27-JUN-2003; 2003US-0482719P.
XX
PΑ
     (UYTE-) UNIV TEL AVIV FUTURE TECHNOLOGY DEV LP.
XX
PΙ
     Eldar-Finkelman H;
XX
DR
     WPI; 2004-468713/44.
XX
PT
     Conjugate useful for treating biological condition e.g., obesity, insulin
PT
     -dependent condition, neurodegenerative disease or psychotic disease,
PT
     comprises polypeptide having hydrophobic moiety that inhibits glycogen
PT
     snythase kinase-3.
XX
PS
     Example; SEQ ID NO 10; 95pp; English.
XX
CC
     The invention relates to a conjugate (I) comprising a specific
CC
     polypeptide and one or more hydrophobic moieties attached to the
     polypeptide, where (I) is capable of inhibiting an activity of glycogen
CC
CC
     synthase kinase-3 (GSK-3). Further disclosed is a pharmaceutical
CC
     composition (II) comprising (I) as an active ingredient. Conjugates of
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CC
     signaling and Up-regulators of beta-catenin. The conjugate of the
CC
     invention is useful for treating a biological condition chosen from
CC
     obesity, non-insulin dependent diabetes mellitus, an insulin-dependent
     condition, an affective disorder, a neurodegenerative disease or disorder
CC
     and a psychotic disease or disorder. The affective disorder is chosen
CC
     from a unipolar disorder and a bipolar disorder. The unipolar disorder is
CC
CC
     depression. The bipolar disorder is manic depression. The
CC
     neurodegenerative disorder results from an event chosen from cerebral
CC
     ischaemia, stroke traumatic brain injury and bacterial infection. The
     neurodegenerative disorder is a chronic neurodegenerative disorder. The
CC
CC
     chronic neurodegenerative disorder results from a disease chosen from
CC
     Alzheimer's disease, Huntington's disease, Parkinson's disease, AIDS
CC
     associated dementia, amyotrophic lateral sclerosis (AML) and multiple
CC
     sclerosis. The psychotic disorder is schizophrenia. The pharmaceutical
CC
     composition is useful for treating biological conditions as mentioned
CC
     above. The current sequence represents a GSK-3 inhibitor peptide.
XX
SO
     Sequence 11 AA;
  Query Match
                          100.0%; Score 63; DB 8; Length 11;
  Best Local Similarity
                          100.0%; Pred. No. 0.13;
  Matches 11; Conservative 0; Mismatches
                                                   0; Indels
                                                                 0; Gaps
            1 KEEPPAPPOSP 11
Qу
              1 KEEPPAPPQSP 11
Db
RESULT 5
ADA54166
ID
     ADA54166 standard; protein; 305 AA.
XX
AC
     ADA54166;
XX
DT
     20-NOV-2003 (first entry)
XX
DE
     Human protein, SEQ ID 1734.
XX
     Cytostatic; Anti-inflammatory; Osteopathic; Neuroprotective; Nootropic;
KW
KW
     Gene Therapy; human; secretory protein; membrane proteins; cancer;
     inflammatory disease; osteoporosis; neurological disease.
KW
XX
OS
     Homo sapiens.
XX
PN
     EP1293569-A2.
XX
PD
     19-MAR-2003.
XX
PF
     21-MAR-2002; 2002EP-00006586.
XX
PR
     14-SEP-2001; 2001JP-00328381.
PR
     24-JAN-2002; 2002US-0350435P.
XX -
PΑ
     (HELI-) HELIX RES INST.
PΑ
     (REAS-) RES ASSOC BIOTECHNOLOGY.
```

the invention act as inhibitors of GSK-3, stimulators of insulin

CC

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XX
      Isogai T, Sugiyama T, Otsuki T, Wakamatsu A, Sato H, Ishii S;
 PΙ
      Yamamoto J, Isono Y, Hio Y, Otsuka K, Nagai K, Irie R, Tamechika I;
 ΡI
      Seki N, Yoshikawa T, Otsuka M, Nagahari K, Masuho Y;
 ΡI
 XX
 DR
      WPI; 2003-395539/38.
      N-PSDB; ADA52527.
 DR
 XX
 PT
      New polynucleotides encoding full-length polypeptides, e.g. secretory
 PT
      and/or membrane proteins, useful for developing medicines for diseases in
 PT
      which the gene is involved, or as target molecules for gene therapy.
 XX
 PS
      Claim 14; SEQ ID NO 1734; 205pp; English.
 XX
 CC
      The present invention relates to novel human secretory or membrane
 CC
      proteins (ADA54072-ADA55710) and their coding sequences (ADA52433-
      ADA54071). The coding sequences are useful in the gene therapy of
 CC
 CC
      diseases caused by abnormalities of the proteins, e.g. cancer,
 CC
      inflammatory diseases, osteoporosis or neurological disease.
 XX
 SQ
      Sequence 305 AA;
   Query Match
                           95.2%; Score 60; DB 6; Length 305;
   Best Local Similarity
                           90.9%; Pred. No. 6.9;
   Matches
             10; Conservative
                                  1; Mismatches
                                                    0; Indels
                                                                  0;
                                                                      Gaps
                                                                              0;
             1 KEEPPAPPOSP 11
. Qy
               ||||||
 Db
            .87 KEEPPSPPQSP 97
 Search completed: November 3, 2005, 09:07:08
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Job time : 179 secs